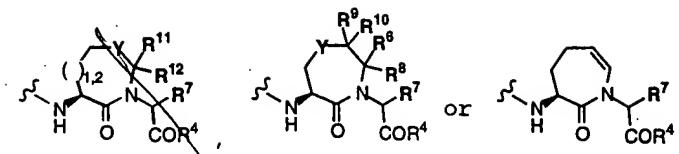


Cont



where X is CH_2 ,

R⁷, R⁸ and R⁹ are independently selected from hydrogen, alkyl, alkenyl, cycloalkyl-(CH_2)_m-, aryl-(CH_2)_m- and heteroaryl-(CH_2)_m-;

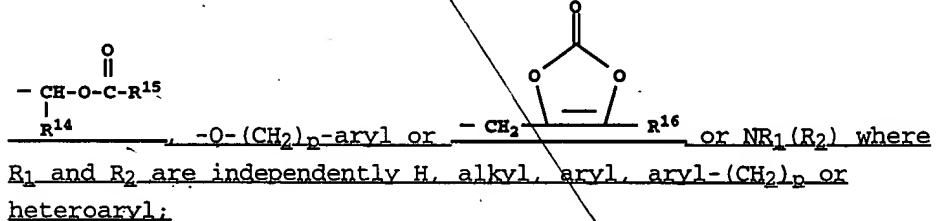
where m is 0 or an integer from 1 to 6;

R⁶, R¹⁰, R¹¹, and R¹² are independently selected from hydrogen, alkyl, alkenyl, cycloalkyl-(CH_2)_p-, aryl-(CH_2)_p- and heteroaryl-(CH_2)_p-; and

R⁴ is OH, Oalkyl, O-(CH_2)_p-heteroaryl.

A1

Cont



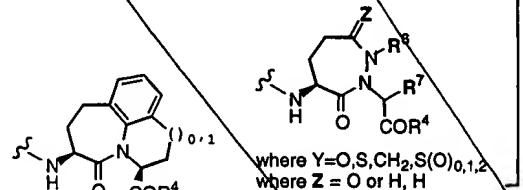
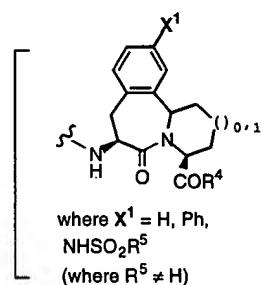
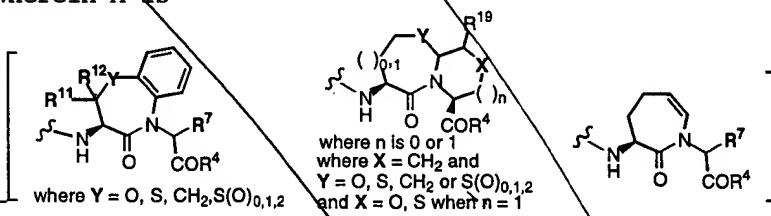
R¹⁴ is hydrogen, alkyl, cycloalkyl, or phenyl;

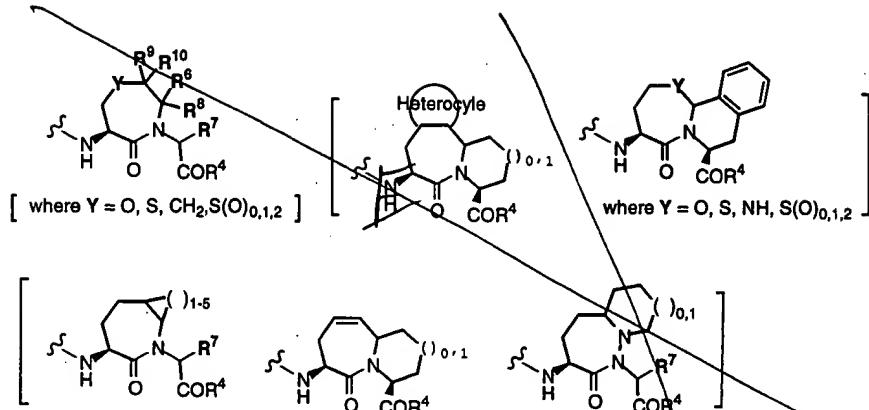
R¹⁵ is hydrogen, alkyl, alkoxy or phenyl; and

R¹⁶ is alkyl or aryl-(CH_2)_m-.

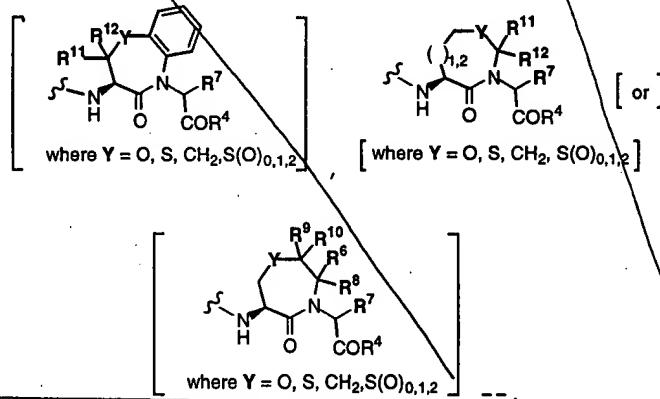
1-6. (Amended) The compound as defined in Claim 1
wherein A is

Board Reason





--7. (Amended) The compound as defined in Claim [6] 1
wherein A is



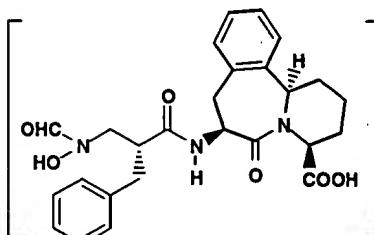
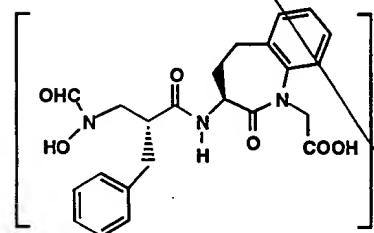
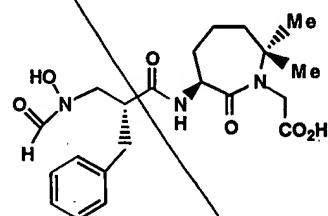
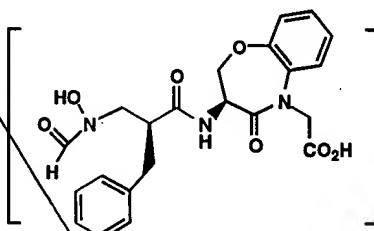
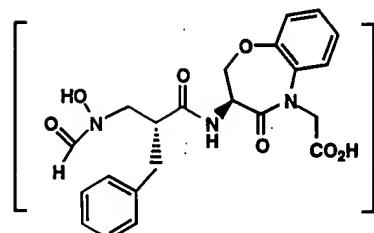
B
X
Sub
E1
Board Decision

--12. (Amended) A pharmaceutical composition comprising a [therapeutically effective amount of a] compound as defined in Claim 1 and a pharmaceutically acceptable carrier therefor.--

--13. (Amended) The pharmaceutical composition as defined in Claim 12 useful in the treatment of [cardiovascular diseases such as] hypertension and/or congestive heart failure.--

--14. (Amended) A method of treating [a cardiovascular disease such as] hypertension and/or congestive heart failure, which comprises administering to a mammalian species a therapeutically effective amount of a composition as defined in Claim 12.--

--15. (Amended) The compound as defined in Claim 1 which is



or a pharmaceutically acceptable salt thereof. --